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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-13-12MW]

Proposed Data Collections Submitted for
Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-7570 or send comments to Kimberly S. Lane, at 1600 Clifton Road, MS D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

## Proposed Project

Hepatitis Testing and Linkage to Care Monitoring & Evaluation System - New-National Center for HIV/AIDS, Viral Hepatitis, STD and TB

Prevention (NCHHSTP), Centers for Disease Control and Prevention

(CDC).

## Background and Brief Description

The National Center for HIV/AIDS, Viral Hepatitis, STD, and TB

Prevention (NCHHSTP), Centers for Disease Control and Prevention is

requesting a three-year OMB approval for establishing a Hepatitis

Testing and Linkage to Care (HEPTLC) Monitoring and Evaluation System

to collect standardized, non-identifying, client-level and test-level

hepatitis testing information from funded testing sites at multiple

settings. Grantees will be required to use this web-based HEPTLC

software application to collect and report testing and linkage to care

activities.

The HEPTLC data collection and reporting system will enable CDC to receive standardized, non-identifying information from funded grantees, including: 1) Information about test sites that provide HEPTLC services and laboratories that provide lab testing; 2) Information about testing participants, including demographics, risk characteristics, vaccination history, etc.

3) Information related to diagnostic test results; and 4) Information about post-test follow-ups, including notification of test result, post-test-counseling, linkage to care and preventive services, and case report to surveillance authorities. CDC will use HEPTLC data for

the following purposes: (1) Monitor the implementation activities of the HEPTLC initiative, as well as evaluate the progress and performance made by the grantees. Findings will further inform strategic planning and program improvement; (2) Inform recommendations and strategies of increasing early identification of infected persons and linkage to care, based on participant characteristics and linkage to care among those persons who are infected; (3) Identify best practices and gaps in implementing HEPTLC in various testing settings, and quide CDC in providing technical assistance to the grantees; (4) Produce standardized and specialized reports that will inform grantees, CDC Project Officers, HHS, Congress and other stakeholders of the process, outcome and accountability measures; (5) Assess public health prevention funds and resources allocations with respect to prioritized risk populations; (6) Advocate the needs for priority setting and budget allocation for hepatitis prevention. Funded sites will use HEPTLC data for the following purposes: (1) Understand targeted populations (demographics, risk behaviors, vaccination histories, etc) and assess the extent to which the targeted populations have been reached; (2) Document how well the project is progressing in meeting goals/objectives set forth by CDC (e.g. who delivered what to whom, how many, where, when, and how well), as well as performance indicators related to testing, counseling and linkage to care; (3) Highlight opportunities for local program collaboration and service integration (PCSI) to prevent hepatitis: (4) Fulfill data collection and reporting requirements

outlined in the cooperative agreements.

The data will enable CDC to be accountable for the funding it provides, the populations that are served, the services being provided, and for the strategies and practices effectiveness in implementing HEPTLC. The data will also enable CDC to be accountable to the administration, Congress, or other stakeholders for the proper use of public money or provide transparency for the programs it funds. Respondents will be testing sites at multiple settings, including health departments, community based organizations (CBOs), community health centers (CHCs), person who inject drugs (PWID) treatment centers, and other settings, e. g. HIV or STD clinics, Federally Qualified Health Centers (FQHCs). They will routinely collect, enter, and report information about the test site, client demographics and behaviors, testing results and linkage to care follow up information within the web-based HEPTLC system. CDC anticipates that routine information collection will begin once OMB approval is received and will be carried out through the project period September 2012 -September 2013.

There are no costs to respondents other than their time. The total estimated annual burden hours are 6000.

## Estimated Annualized Burden Hours

Type of Respondents	Form Name	Number of Respondents	Responses per Respondent	Average Burden per response (in hours)
<pre>HBV - CBOs/Health Jurisdictions HCV - multiple sites (IDU, CHCs, Others, ECHO)</pre>	HEPTLC Data Variables & Values (test- level monthly reporting)	40	12	12
<pre>HBV - CBOs/Health Jurisdictions HCV - multiple sites (IDU, CHCs,</pre>	HEPTLC Template (program- level	40	4	1.5

Type of Respondents	Form Name	Number of Respondents	Responses per Respondent	Average Burden per response (in hours)
Others, ECHO)	reporting/qua rterly)			

Date: October 22, 2012

Ron A. Otten, Ph.D.

Director, Office of Scientific Integrity (OSI)
Office of the Associate Director for Science
(OADS)

Office of the Director

Centers for Disease Control and Prevention

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